Amendments to the Claims:

This listing of claims will replace all prior versions and listings of claims in the specification.

Listing of Claims:

Claim 1 (Original): A pharmaceutical composition comprising:

(a) an anticholinergic selected from glycopyrronium bromide or an ester of a bi- or tricyclic amino alcohol of formula (I)

$$z$$
 O Q N^{+} $R'_{(I)}$

wherein:

Q is one of the groups -CH₂-CH₂-, -CH=CH-, or

R is methyl, ethyl, or propyl optionally substituted by fluorine or hydroxy,

R' is methyl, ethyl, or propyl, and

an equivalent of an anion X counters the positive charge of the N atom; and

Z is one of the groups

$$R^1$$
 or R_1

wherein:

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Y is a single bond or an O atom,

R¹ is hydrogen, hydroxy, methoxy, ethoxy, propoxy, methyl, ethyl, propyl, hydroxymethyl, hydroxyethyl, or hydroxypropyl,

R² is a thienyl, phenyl, or cyclohexyl group, wherein these groups are optionally substituted by methyl, and thienyl and phenyl are optionally substituted by fluorine or chlorine, and

R³ is hydrogen, or a thienyl or phenyl group which is optionally substituted by fluorine, chlorine, or methyl; and

(b) a betamimetic selected from the group consisting of: formoterol; salmeterol; 4-hydroxy-7-[2-{[2-{[3-(2-phenylethoxy)propyl]sulfonyl}ethyl]amino}ethyl]-2(3H)-benzothiazolone; 1-(2-fluoro-4-hydroxyphenyl)-2-[4-(1-benzimidazolyl)-2-methyl-2-butylamino]ethanol; 1-[3-(4-methoxybenzylamino)-4-hydroxyphenyl]-2-[4-(1-benzimidazolyl)-2-methyl-2-1-[2*H*-5-hydroxy-3-oxo-4*H*-1,4-benzoxazin-8-yl]-2-[3-(4-*N*,*N*butylamino]ethanol; dimethylaminophenyl)-2-methyl-2-propylaminolethanol; 1-[2H-5-hydroxy-3-oxo-4H-1,4benzoxazin-8-yl]-2-[3-(4-methoxyphenyl)-2-methyl-2-propylamino]ethanol; 1-[2*H*-5hydroxy-3-oxo-4*H*-1,4-benzoxazin-8-yl]-2-[3-(4-*n*-butyloxyphenyl)-2-methyl-2-1-[2H-5-hydroxy-3-oxo-4H-1,4-benzoxazin-8-yl]-2-{4-[3-(4propylaminolethanol; and methoxyphenyl)-1,2,4-triazol-3-yl]-2-methyl-2-butylamino}ethanol, and pharmacologically compatible acid addition salt thereof.

Claim 2 (Original): The pharmaceutical composition according to claim 1, wherein the anticholinergic is an ester of a bi- and tricyclic amino alcohol of formula (I)

$$z$$
 O
 Q
 $R'_{(I)}$

wherein:

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Q is one of the groups -CH₂-CH₂-, -CH=CH-, or

R is methyl or ethyl,

R' is methyl, and

anion X is bromide; and

Z is one of the groups

$$R^1$$
 or R_1

wherein:

R¹ is hydrogen, hydroxy, or hydroxymethyl,

R² is a thienyl, phenyl, or cyclohexyl group, and

R³ is hydrogen, or a thienyl or phenyl group.

Claim 3 (Original): The pharmaceutical composition according to claim 1, wherein the anticholinergic is a salt of tiotropium.

Claim 4 (Original): The pharmaceutical composition according to claim 1, wherein the anticholinergic is tiotropium bromide.

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Claim 5 (Original): The pharmaceutical composition according to claim 1, wherein the

betamimetic is formoterol or salmeterol, or a pharmacologically compatible acid addition salt

thereof

Claim 6 (Original): The pharmaceutical composition according to claim 1, wherein the

anticholinergic is tiotropium bromide and the betamimetic is formoterol, or a pharmacologically

compatible acid addition salt thereof.

Claim 7 (Original): The pharmaceutical composition according to claim 1, wherein the

anticholinergic is tiotropium bromide and the betamimetic is salmeterol, or a pharmacologically

compatible acid addition salt thereof.

Claim 8 (Original): The pharmaceutical composition according to claim 1, wherein the anion X

is selected from the group consisting of: chloride, bromide, and methanesulfonate,

Claim 9 (Original): The pharmaceutical composition according to one of claims 1 to 8, wherein

the pharmaceutical composition is an inhaled pharmaceutical composition.

Claim 10 (Original): A process for the production of a pharmaceutical composition according to

one of claims 1 to 8, comprising:

(a) mixing the anticholinergic and the betamimetic; and optionally

(b) adding an adjuvant and/or carrier materials.

Claims 11-14 (Cancelled).

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